



Day : Friday
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MATRIXES	13214
((BONE ADJ MARROW) ADJ (EXTRACELLULAR ADJ MATRIX)).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	27
((((BONE ADJ MARROW) ADJ (EXTRACELLULAR ADJ MATRIX))).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	27

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OP=AND

<u>L18</u>	((bone adj marrow) adj (extracellular adj matrix))	27	<u>L18</u>
<u>L17</u>	L16 and (tissue adj (repair or engineering))	82	<u>L17</u>
<u>L16</u>	L3 and (transgenic)	180	<u>L16</u>
<u>L15</u>	(stimulus) same ((in adj vivo) and matrix)	10	<u>L15</u>
<u>L14</u>	L13 not L5	0	<u>L14</u>
<u>L13</u>	L12 and L3	29	<u>L13</u>
<u>L12</u>	L4 same (donor)	1321	<u>L12</u>
<u>L11</u>	L5 not L6	105	<u>L11</u>
<u>L10</u>	L6 not L7	76	<u>L10</u>
<u>L9</u>	L7 not L8	56	<u>L9</u>
<u>L8</u>	L7 and (VEGF)	51	<u>L8</u>
<u>L7</u>	L6 and (bone adj marrow)	107	<u>L7</u>
<u>L6</u>	L5 and (vector or transfected)	183	<u>L6</u>
<u>L5</u>	L4 and L3	288	<u>L5</u>
<u>L4</u>	(Preconditioning or conditioning or preconditioned or conditioned or stimulus) same ((in adj vivo) or tissue or organ or cell)	54696	<u>L4</u>
<u>L3</u>	(Decellularized or acellular or decellularisation or decellularization) same (tissue or matrix or matrices)	1405	<u>L3</u>
<u>L2</u>	L1 and ((decellularized adj extracellular) adj matrix)	3	<u>L2</u>
<u>L1</u>	Freyman-Toby.in.	51	<u>L1</u>

END OF SEARCH HISTORY

Welcome to DialogClassic Web(tm)

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Last logoff: 26oct07 11:56:11
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*** ANNOUNCEMENTS ***

NEW FILES RELEASED

***BIOSIS Previews Archive (File 552)
***BIOSIS Previews 1969-2007 (File 525)
***Trademarkscan - South Korea (File 655)

RESUMED UPDATING

***File 141, Reader's Guide Abstracts

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***File 5, BIOSIS Previews - archival data added
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NEWS

Chemical Structure Searching now available in Prous Science Drug Data Report (F452), Prous Science Drugs of the Future (F453), IMS R&D Focus (F445/955), Pharmaprojects (F128/928), Beilstein Facts (F390), Derwent Chemistry Resource (F355) and Index Chemicus (File 302).

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Cost is in DialUnits

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B 155, 5, 73

02nov07 14:53:10 User259876 Session D1048.1

\$0.98 0.279 DialUnits File1

\$0.98 Estimated cost File1

\$0.05 INTERNET

\$1.03 Estimated cost this search

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File 155:MEDLINE(R) 1950-2007/Oct 31

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3264403 TISSUE
481080 MATRIX
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1339881 VIVO
1294235 IN(W) VIVO
3264403 TISSUE
545161 ORGAN
5781231 CELLS
313594 DONOR
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S S1 AND S2
3991 S1
137086 S2
S3 76 S1 AND S2
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76 S3
347610 VECTOR
170190 TRANSFECTED
S4 0 S3 AND (VECTOR OR TRANSFECTED)
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S5 39 RD S3 (unique items)
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S S5 NOT PY>2003
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S6 19 S5 NOT PY>2003

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S S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
19 S6
3264403 TISSUE
359561 REPAIR
325834 ENGINEERING
37197 TISSUE(W) (REPAIR OR ENGINEERING)
S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))

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T S7/3,K/ALL

7/3,K/1 (Item 1 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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14003792 PMID: 12415348

Experimental abdominal wall defect repaired with acellular matrix.
Gamba P G; Conconi M T; Lo Piccolo R; Zara G; Spinazzi R; Parnigotto P P
Department of Pediatric Surgery, University of Padua, Italy. g
piergioorgio@hotmail.com
Pediatric surgery international (Germany) Sep 2002, 18 (5-6) p327-31
, ISSN 0179-0358--Print Journal Code: 8609169
Publishing Model Print-Electronic
Document type: Journal Article
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Experimental abdominal wall defect repaired with acellular matrix .
... of a non-immunogenic and non-prosthetic biomaterial that could guide
the regeneration of normal tissue is a fascinating possibility.
Biomaterials are already in use, but in our experience, an acellular
matrix (ACM) can stimulate exact regeneration of the absent tissue . We
explored the possibility of using an ACM to repair a muscular AWD in an...

... oblique muscle was resected (3 x 3 cm). The animals underwent
reconstruction with homologous diaphragm acellular matrix (HDAM) grafts
that were previously prepared using a detergent enzymatic method. The
patches were evaluated...

... each group; moreover, 90 days post-surgery an electromyogram (EMG) (n =
6) of the implanted matrix was recorded. Histologic analysis demonstrated
that the HDAM supported fibroblast migration, deposition of newly-formed...

... not able to produce reconstruction of the skeletal muscle, and was
progressively remodeled into fibrous tissue . Since the ultimate reason
for failure of muscle regeneration is a lack of myogenesis, future studies
will use ACMS preconditioned by various regulators of myoblast
proliferation and differentiation.

Descriptors: *Abdominal Wall--abnormalities--AB; *Abdominal Wall
--surgery--SU; *Biocompatible Materials; * Tissue Engineering
?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECEL-

LULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
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 ITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR
 CELLS OR DONOR)
 S3 76 S1 AND S2
 S4 0 S3 AND (VECTOR OR TRANSFECTED)
 S5 39 RD S3 (unique items)
 S6 19 S5 NOT PY>2003
 S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
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T S6/3,K/ALL

6/3,K/1 (Item 1 from file: 155)
 DIALOG(R)File 155:MEDLINE(R)
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24212957 PMID: 17177738

Development of a multilayered in vitro model for studying events
 associated with wound healing.

Stephens P; Wood E J; Raxworthy M J

Department of Oral Surgery, Medicine and Pathology, Dental School,
 University of Wales College of Medicine, Cardiff, Wales, UK.

Wound repair and regeneration - official publication of the Wound Healing
 Society and the European Tissue Repair Society (United States) Jul-Sep
 1996, 4 (3) p393-401, ISSN 1067-1927--Print Journal Code: 9310939

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: PubMed not MEDLINE

...the study of cutaneous wound repair processes. This has stimulated the
 investigation of three-dimensional tissue equivalent systems such as the
 dermal and skin equivalent models. With the use of a...

... wounded dermal equivalent (bilayered model) or skin equivalent
 (tri-layered model) was placed onto an acellular collagen lattice and
 fixed in place with polymerizing collagen. This model permitted observation
 of the...

...number of fibroblasts in this space increased dramatically over a period
 of 9 days, the cells appearing to migrate both over and through the
 acellular lower collagen layer. Keratinocyte reepithelialization of the
 "wound space" was completed after 5 days. With...

... model described here should facilitate the study of fibroblast and
 keratinocyte responses to a wound stimulus in vitro and be a plausible in
 vitro system for evaluating agents which may have...

6/3,K/2 (Item 2 from file: 155)
 DIALOG(R)File 155:MEDLINE(R)
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14109190 PMID: 12641817

An in vitro examination of an extracellular matrix scaffold for use in
 wound healing.

Solomon Denis E

Clinical Research Division, Department of Surgery, University of Miami,

School of Medicine, Miami, Florida 33101, USA. denissolomon@yahoo.com
 International journal of experimental pathology (England) Oct 2002, 83
 (5) p209-16, ISSN 0959-9673--Print Journal Code: 9014042
 Publishing Model Print
 Document type: Journal Article
 Languages: ENGLISH
 Main Citation Owner: NLM
 Record type: MEDLINE; Completed

This paper describes evidence that an extracellular matrix (ECM) secreted by human umbilical vein endothelial cells (HUVECs) assembled on gelatin coated plates overlaid by a mixed matrix secreted by human dermal microvascular endothelial cells (HDMECs) and human dermal fibroblasts provides a viable acellular scaffold for use in wound healing. Trypsinized epidermal keratinocytes or colonies from Dispase-digested fresh and cadaver skin tissue adhered and proliferated on either HUVECs ECM/gelatin or mixed matrix overlaid on HUVECs ECM/gelatin. An epithelial-mesenchymal interaction, previously thought to be tissue-specific, was exposed as well as concomitant integrin versatility. Furthermore, heterologous HDMECs and dermal fibroblasts attached and proliferated on the mixed matrix as well as HUVECs ECM. The conditioned medium from HUVECs (HUVECs CM) was found to neutralize the lingering after effects of Dispase, and could be used for the tissue culture of epidermal keratinocytes, HDMECs and dermal fibroblasts, which share related extracellular secretions. Taken together...

... epithelial autografts can be redesigned to include both epithelial and dermal elements, and advances the acellular 'sandwich' ECM scaffold as a possible structural replacement for the lamina densa and lamina lucida...

6/3,K/3 (Item 3 from file: 155)
 DIALOG(R) File 155:MEDLINE(R)
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14003792 PMID: 12415348

Experimental abdominal wall defect repaired with acellular matrix.
 Gamba P G; Conconi M T; Lo Piccolo R; Zara G; Spinazzi R; Parnigotto P P
 Department of Pediatric Surgery, University of Padua, Italy. g
 piergiorgio@hotmail.com
 Pediatric surgery international (Germany) Sep 2002, 18 (5-6) p327-31
 , ISSN 0179-0358--Print Journal Code: 8609169
 Publishing Model Print-Electronic
 Document type: Journal Article
 Languages: ENGLISH
 Main Citation Owner: NLM
 Record type: MEDLINE; Completed

Experimental abdominal wall defect repaired with acellular matrix .
 ... of a non-immunogenic and non-prosthetic biomaterial that could guide the regeneration of normal tissue is a fascinating possibility. Biomaterials are already in use, but in our experience, an acellular matrix (ACM) can stimulate exact regeneration of the absent tissue . We explored the possibility of using an ACM to repair a muscular AWD in an...

... oblique muscle was resected (3 x 3 cm). The animals underwent reconstruction with homologous diaphragm acellular matrix (HDAM) grafts that were previously prepared using a detergent enzymatic method. The patches were evaluated...

... each group; moreover, 90 days post-surgery an electromyogram (EMG) (n = 6) of the implanted matrix was recorded. Histologic analysis demonstrated that the HDAM supported fibroblast migration, deposition of newly-formed...

... not able to produce reconstruction of the skeletal muscle, and was progressively remodeled into fibrous tissue. Since the ultimate reason for failure of muscle regeneration is a lack of myogenesis, future studies will use ACMS preconditioned by various regulators of myoblast proliferation and differentiation.

6/3,K/4 (Item 4 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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13943414 PMID: 12296587

Bone repair following recombinant human bone morphogenetic protein-2 stimulated periodontal regeneration.

Selvig Knut A; Sorensen Rachel G; Wozney John M; Wikesjo Ulf M E
Department of Dental Research, University of Bergen, Faculty of Dentistry, Norway. knut.selvig@odont.uib.no

Journal of periodontology (United States) Sep 2002, 73 (9) p1020-9,
ISSN 0022-3492--Print Journal Code: 8000345

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... therapy for periodontal regeneration. The objective of this study was to characterize, in some detail, tissue reactions following surgical implantation of rhBMP-2/ACS into periodontal defects. METHODS: Four young adult...

... implantation of rhBMP-2/ACS into large supra-alveolar periodontal defects resulted in a variable tissue response without marked difference between 4- and 8-week observations. New bone, exceeding the volume of the normal alveolar process, had formed within 4 weeks. The regenerated bone tissue consisted of finely trabeculated woven bone. Marrow spaces exhibited a continuous lining of osteoblasts, osteoclasts, and resting cells. The marrow spaces contained numerous large, thin-walled vessels but were almost devoid of collagen...

... were free of structured elements except for occasional aggregates of effete erythrocytes. A variety of tissue reactions were observed along the root surface including areas of resorption, areas of hard tissue deposition, and areas without resorptive or appositional activity. Ankylosis was a frequent observation, although areas showing characteristics of a periodontal ligament with a fine layer of acellular fiber cementum and occasional inserting Sharpey's fibers were also observed. Osteoblasts facing the root...

... along and onto the instrumented adjacent root surface. Lamellated trabecular bone was the predominant regenerated tissue. A typical cementum-periodontal ligament-alveolar bone relationship was a rare observation. The great variability in histological tissue response along the instrumented root surface indicates that the stimulus to hard tissue formation resided primarily in the rhBMP-2/ACS implant rather than in the root surface.

6/3,K/5 (Item 5 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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13220634 PMID: 11350647

Keratinocyte-driven contraction of reconstructed human skin.

Chakrabarty K H; Heaton M; Dalley A J; Dawson R A; Freedlander E; Khaw P T; Mac Neil S

Section of Medicine, Division of Clinical Sciences and Plastics, Burns and Reconstructive Surgery, Northern General Hospital NHS Trust, Sheffield and Institute of Ophthalmology and Moorfields Eye Hospital, London, United Kingdom.

Wound repair and regeneration - official publication of the Wound Healing Society and the European Tissue Repair Society (United States) Mar-Apr 2001, 9 (2) p95-106, ISSN 1067-1927--Print Journal Code: 9310939

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

We have previously reported that reconstructed human skin, using deepidermized acellular sterilized dermis and allogeneic keratinocytes and fibroblasts, significantly contracts in vitro. Contracture of split skin...

... several approaches to prevent or reduce contraction. Three different methodologies for sterilization of the dermal matrix were examined: glycerol, ethylene oxide and a combination of glycerol and ethylene oxide. While the nature of the sterilization technique influenced the extent of contraction and thinner dermal matrices contracted proportionately more than thicker matrices, in all cases contraction was driven by the keratinocytes with relatively little influence from the fibroblasts. The contraction of the underlying dermis did not represent any change in tissue mass but rather a reorganization of the dermis which was rapidly reversed (within minutes) when...

... phosphate to be ineffective and ascorbic acid-2-phosphate to exacerbate contraction. However, Galardin, a matrix metalloproteinase inhibitor and keratinocyte conditioned media, both inhibited contraction.

6/3,K/6 (Item 6 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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12850538 PMID: 10972644

Porcine fetal enamel matrix derivative stimulates proliferation but not differentiation of pre-osteoblastic 2T9 cells, inhibits proliferation and stimulates differentiation of osteoblast-like MG63 cells, and increases proliferation and differentiation of normal human osteoblast NHOst cells.

Schwartz Z; Carnes D L; Pulliam R; Lohmann C H; Sylvia V L; Liu Y; Dean D D; Cochran D L; Boyan B D

Department of Orthopaedics, University of Texas Health Science Center, San Antonio 78229-3900, USA.

Journal of periodontology (UNITED STATES) Aug 2000, 71 (8) p1287-96, ISSN 0022-3492--Print Journal Code: 8000345

Contract/Grant No.: DE-08603; DE; NIDCR

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't;

Research Support, U.S. Gov't, P.H.S.

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

BACKGROUND: Embryonic enamel matrix proteins are hypothesized to be involved in the formation of acellular cementum during tooth development, suggesting that these proteins can be used to regenerate periodontal tissues. Enamel matrix protein derived from embryonic porcine tooth germs is used clinically, but the mechanisms by which...

... examined the response of osteoblasts at 3 stages of osteogenic maturation to porcine fetal enamel matrix derivative (EMD). Proliferation (cell number and [3H]-thymidine incorporation), differentiation (alkaline phosphatase and osteocalcin), matrix synthesis ([35S]-sulfate incorporation; percentage of collagen production), and local factor production (prostaglandin E2 [PGE2] and transforming growth factor-beta 1 [TGF-beta1]) were measured in cultures of 2T9 cells (pre-osteoblasts which exhibit osteogenesis in response to bone morphogenetic protein-2 [BMP-2]), MG63 human osteoblast-like osteosarcoma cells, and normal human osteoblasts (NHOst cells). RESULTS: EMD regulated osteoblast proliferation and differentiation, but the effects were cell-specific. In 2T9...

... proliferation but had no effect on alkaline phosphatase-specific activity. EMD decreased proliferation of MG63 cells and increased cellular alkaline phosphatase and osteocalcin production. There was no effect on collagen synthesis, proteoglycan sulfation, or PGE2 production; however, TGF-beta1 content of the conditioned media was increased. There was a 60-fold increase in cell number in third passage NHOst cells cultured for 35 days in the presence of EMD. EMD also caused a biphasic increase...

...day 14. CONCLUSIONS: EMD affects early states of osteoblastic maturation by stimulating proliferation, but as cells mature in the lineage, EMD enhances differentiation.

6/3,K/7 (Item 7 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

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12558276 PMID: 10505815

Subgingival acellular dermal matrix allograft for the treatment of gingival recession: a case report.

Tal H

Department of Periodontology, The Maurice and Gabriela Goldschleger School of Dental Medicine, Tel Aviv University, Israel.

Journal of periodontology (UNITED STATES) Sep 1999, 70 (9) p1118-24, ISSN 0022-3492--Print Journal Code: 8000345

Publishing Model Print

Document type: Case Reports; Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subgingival acellular dermal matrix allograft for the treatment of gingival recession: a case report.

Root coverage procedures using subgingival soft tissue grafts or guided

tissue regeneration have attracted much interest within the past 2 decades. Recently, acellular dermal matrix allograft (ADMA) has been introduced as a substitute for palatal donor tissue in gingival augmentation procedures. This study was undertaken to examine the potential of ADMA to be used as a substitute for autogenous connective tissue graft material in a root coverage procedure in a case with moderate gingival recession combined with reduced keratinized attached gingiva. After thorough root planing and conditioning of the root surface with a saturated solution of tetracycline-HCl, a trapezoidal mucoperiosteal flap ...

... observations, it is suggested that ADMA may be a possible substitute to free autogenous connective tissue grafts and/or bioabsorbable barrier membranes. Further clinical and histological studies are necessary to understand...

6/3,K/8 (Item 8 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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11492761 PMID: 9326143

Keratinocytes contract human dermal extracellular matrix and reduce soluble fibronectin production by fibroblasts in a skin composite model.
Ralston D R; Layton C; Dalley A J; Boyce S G; Freedlander E; MacNeil S
University Department of Medicine, Northern General Hospital, Sheffield, UK.

British journal of plastic surgery (ENGLAND) Sep 1997, 50 (6)
p408-15, ISSN 0007-1226--Print Journal Code: 2984714R
Publishing Model Print
Document type: Journal Article; Research Support, Non-U.S. Gov't
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Composites of human de-epidermised acellular dermis and normal adult human keratinocytes and fibroblasts were examined for the ability of cells to contract these composites. Image analysis of the outline of the composites showed that, in...

... was no significant contraction of the dermis with fibroblasts alone or in the absence of cells. The presence or absence of basement membrane antigens did not influence the effect of keratinocytes on dermal contraction. Analysis of the conditioned media from these composites showed that the greatest fibronectin production was seen with fibroblasts alone...

... the presence and absence of the basement membrane, indicating that keratinocytes modify dermal fibroblast extracellular matrix production. This study shows that while keratinocytes and fibroblasts are clearly influencing each other's...

... is the keratinocyte and not the fibroblast which causes contraction of the human de-epidermised acellular dermis.

6/3,K/9 (Item 9 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

11364892 PMID: 9183827

Silicone breast implants: pathology.

Raso D S; Greene W B

Department of Pathology, Medical University of South Carolina,
Charleston, USA.

Ultrastructural pathology (UNITED STATES) May-Jun 1997, 21 (3)
p263-71, ISSN 0191-3123--Print Journal Code: 8002867

Publishing Model Print

Document type: Journal Article; Review

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... legal system. Pathologists must be aware of the controversy and treat each SBI and associated tissue as a potential lawsuit. Grossly, silicone is a clear, viscous substance that may be observed...

... cases, a fibrous capsule surrounds the SBI, with the interface lining varying from a virtually acellular to a synovial-like lining composed of phagocytic and secretory cells. Silicone can often be identified within the fibrous capsule and also in distant tissues biopsied...

... without ultrastructural evidence of cytologic effects. This study has demonstrated that silicone accumulates at distant tissue sites due to preexisting inflammation acting as a stimulus. Thus, silicone is not a primary inducer of inflammatory disease processes. These findings are supported...

6/3,K/10 (Item 10 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

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10329146 PMID: 7890329

Blood monocytes and spleen macrophages differentiate into microglia-like cells on monolayers of astrocytes: morphology.

Sievers J; Parwaresch R; Wottge H U

Department of Anatomy, University of Kiel, FRG.

Glia (UNITED STATES) Dec 1994, 12 (4) p245-58, ISSN 0894-1491--
Print Journal Code: 8806785

Publishing Model Print

Document type: Comparative Study; Journal Article; Research Support,
Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Several morphological and functional properties of microglial cells, the resident immunoeffector cells of the central nervous system (CNS), differ from those of monocytes/macrophages in other tissues...

... tested the hypothesis that some morphological and functional properties of microglia are induced in myelomonocytic cells by nervous tissue, specifically astrocytes. In the present in vitro studies we compared the differentiation of microglia, blood monocytes, and spleen macrophages on acellular substrates and on monolayers of astrocytes and fibroblasts. On acellular substrates, microglial cells at first acquire an ameboid morphology; later they show a few short, unbranched processes. On monolayers of pure astrocytes, microglial cells at first also differentiate into ameboid cells, but after 5 to 7 days they start to develop processes with large lamellopodial tips...

... territory around the small ellipsoid cell body. By contrast, on monolayers of fibroblasts the microglial cells develop an ameboid morphology, but do not grow the typical long branched processes of the ramified form. Blood monocytes and spleen macrophages behave indistinguishably from microglia both on acellular and cellular substrates, i.e., on astroglia they develop the ramified form, while on fibroblasts...

...of astrocytic monolayers, i.e., physically separated from the astroglia, but exposed to the medium conditioned by astrocytes, a significant proportion of them also develop the ramified shape. These findings indicate ...

... monocytes and macrophages, we take this to be further evidence for the proposition that microglial cells are derived from the myelomonocytic lineage, and, moreover, that properties of resident macrophages are largely determined by tissue components of their host organ .

6/3,K/11 (Item 11 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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09073314 PMID: 1721917

Combined effects of extracellular matrix and growth factors on NBT-II rat bladder carcinoma cell dispersion.
Tucker G C; Boyer B; Valles A M; Thiery J P
Laboratoire de Physiopathologie du Developpement, CNRS URA 1337 Ecole Normale Supérieure, Paris, France.
Journal of cell science (ENGLAND) Oct 1991, 100 (Pt 2) p371-80,
ISSN 0021-9533--Print Journal Code: 0052457
Contract/Grant No.: 1R01 CA 49417-01A2; CA; NCI
Publishing Model Print
Document type: Journal Article; Research Support, Non-U.S. Gov't;
Research Support, U.S. Gov't, P.H.S.
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

... factor and transforming growth factor alpha also promoted NBT-II cell dispersion on glass or tissue culture plastic. We have now further analysed the scatter response to these two growth factors in the presence of extracellular matrix molecules. In the presence of growth factors, no peripheral single-cell dispersion occurred on fibronectin...

...motility inside the monolayer forming around NBT-II aggregates. Patterns of strings or files of cells protruding from the monolayer were often observed. The presence of a scattering activity in the complex acellular extracellular matrix deposited by NBT-II cells themselves strongly suggested that substratum conditioning was responsible for this effect. On the other hand, the two growth factors accelerated collagen...

... to-substratum contact. On laminin or fibronectin and in the presence of growth factors, peripheral cells inside the halo of NBT-II aggregates did not exhibit desmosome linkages. These observations suggest that scatter effects per se are dependent on the composition of the extracellular matrix . In particular, on a substratum nonpermissive for direct cell translocation, individual cell dispersion can be replaced by en bloc patterns of migration following substratum conditioning by the cells .

6/3,K/12 (Item 12 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

08824109 PMID: 1849087

An improved noninfectious murine skin model of organized granulomatous inflammation.

Iida T; Nozaki Y; Fukuyama K; Epstein W L
Department of Dermatology, University of California San Francisco
94143-0536.

Experientia (SWITZERLAND) Mar 15 1991, 47 (3) p273-7, ISSN
0014-4754--Print Journal Code: 0376547

Contract/Grant No.: AR31853; AR; NIAMS

Publishing Model Print

Document type: Journal Article; Research Support, U.S. Gov't, P.H.S.

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... granulomas, originally elicited in naive mice by inoculations of lyophilized hepatic schistosome egg granulomas. The tissue reaction is caused by a single exposure to a noninfectious, acellular granulomagenic stimulus and occurs in healthy mice free of systemic disease. The model should prove useful for...

... analytical dissection of the initiation process. In this study we described the responses of host cells by autoradiography, and light and electron microscopy. The activity of angiotensin-converting enzyme and proline...

6/3,K/13 (Item 13 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

06131504 PMID: 6190818

Capillary endothelial cell cultures: phenotypic modulation by matrix components.

Madri J A; Williams S K

Journal of cell biology (UNITED STATES) Jul 1983, 97 (1) p153-65,
ISSN 0021-9525--Print Journal Code: 0375356

Contract/Grant No.: R01-HL-28373-02; HL; NHLBI

Publishing Model Print

Document type: Journal Article; Research Support, U.S. Gov't, P.H.S.

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Capillary endothelial cells of rat epididymal fat pad were isolated and cultured in media conditioned by bovine aortic endothelial cells and substrata consisting of interstitial or basement membrane collagens. When these cells were grown on interstitial collagens they underwent proliferation, formed a continuous cell layer and, if cultured for long periods of time, formed occasional tubelike structures. In contrast, when these cells were grown on basement membrane collagens, they did not proliferate but did aggregate and form tubelike structures at early culture times. In addition, cells grown on basement membrane substrata expressed more basement membrane constituents as compared with cells grown on interstitial matrices when assayed by immunoperoxidase methods and quantitated by enzyme-linked immunosorbent inhibition assays. Furthermore,

when cells were grown on either side of washed, acellular amnionic membranes their phenotypes were markedly different. On the basement membrane surface they adhered, spread...

... not migrate through the basement membrane. In contrast, when seeded on the stromal surface, these cells were observed to proliferate and migrate into the stromal aspect of the amnion and ultimately formed tubelike structures at high cell densities at longer culture periods (21 d). Thus, connective tissue components play important roles in regulating the phenotypic expression of capillary endothelial cells in vitro, and similar roles of the collagenous components of the extracellular matrix may exist in vivo following injury and during angiogenesis. Furthermore, the culture systems outlined here may be of use in the further study of differentiated, organized capillary endothelial cells in culture.

6/3,K/14 (Item 1 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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17380580 BIOSIS NO.: 200300337323

Annexin II Mediated Plasmin Generation Activates TGFbeta-3 during Epithelial-Mesenchymal Transformation in the Developing Chick Heart.

AUTHOR: Krishnan Suba (Reprint); Deora Arun K (Reprint); Hajjar Katherine A (Reprint)

AUTHOR ADDRESS: Cell and Developmental Biology, Weill Medical College of Cornell University, New York, NY, USA**USA

JOURNAL: Blood 100 (11): pAbstract No. 192 November 16, 2002 2002

MEDIUM: print

CONFERENCE/MEETING: 44th Annual Meeting of the American Society of Hematology Philadelphia, PA, USA December 06-10, 2002; 20021206

SPONSOR: American Society of Hematology

ISSN: 0006-4971

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract

LANGUAGE: English

...ABSTRACT: complexity of these events. Epithelial-mesenchymal transformation (EMT), the process by which subpopulations of endocardial cells of the atrioventricular (AV) region are converted into motile, invasive, mesenchymal cells, is critical to normal AV valve development. The transformed mesenchymal cells invade underlying cardiac jelly and form the fibrous regions of the AV valves of the...

...annexin family of calcium-dependent, peripheral membrane binding proteins, functions as a co-receptor for tissue plasminogen activator (tPA) and plasminogen, enhancing the efficiency of cell-surface generation of the fibrinolytic...

...hypothesize that annexin II, by stimulating localized plasmin generation on the surface of transforming endocardial cells, results in the regulated release of active TGFbeta-3 during AV canal EMT. Serial Northern...

...strong annexin II staining localized specifically to endocardium during AV canal EMT. Primary endocardial endothelial cells (EEC) isolated from embryonic stage 15-18 chick heart explants expressed annexin II protein by...

...an 8-fold increase in the rate of plasmin generation using chick EEC versus an acellular control. Plasmin generation was tPA-, plasminogen-, and cell-surface dependent, and was inhibited apprx90% by...
...of inhibition of transformation to the level of untreated heart explants. Western blot analysis of conditioned medium from heart explant cultures revealed absence of active TGFbeta-3 upon treatment with anti...

...alpha-2 antiplasmin. A TGFbeta bioassay showed a 50% decrease in TGFbeta specific activity in conditioned medium from heart explant cultures treated with anti-annexin II IgG as compared with untreated...

6/3,K/15 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

16667902 BIOSIS NO.: 200200261413

The fibrinolytic receptor, annexin II, mediates epithelial-mesenchymal transformation in the developing avian heart

AUTHOR: Krishnan Suba (Reprint); Deora Arun Kumar B (Reprint); Jacovina Andrew T (Reprint); Lev Emil (Reprint); Hajjar Katherine A (Reprint)

AUTHOR ADDRESS: Pediatrics, Weill Medical College of Cornell University, New York, NY, USA**USA

JOURNAL: Blood 98 (11 Part 1): p788a-789a November 16, 2001 2001

MEDIUM: print

CONFERENCE/MEETING: 43rd Annual Meeting of the American Society of Hematology, Part 1 Orlando, Florida, USA December 07-11, 2001; 20011207

SPONSOR: American Society of Hematology

ISSN: 0006-4971

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: Epithelial-mesenchymal transformation (EMT), the process whereby endocardial cells of the atrioventricular (AV) region of the heart acquire a migratory mesenchymal phenotype, is a critical event in cardiac development. During EMT, endocardial cells remodel extracellular matrix (ECM), invade underlying cardiac jelly, and ultimately form the fibrous portion of the valves and...

...family of calcium-regulated, phospho-lipid binding proteins, that functions as a co-receptor for tissue plasminogen activator (tPA) and plasminogen, and enhances the efficiency of plasmin generation on cell surfaces...

...directly breakdown ECM proteins, or may initiate a proteolytic cascade resulting in activation of pro-matrix metalloproteinases (pro-MMPs). Serine proteases have also been implicated in the activation and liberation of...

...by stimulating localized generation of plasmin on the surface of transforming endocardial or migratory mesenchymal cells of the AV canal. Serial Northern and Western blot analyses of hearts isolated from Hamilton...

...annexin II staining specifically localized to endocardium during ongoing AV canal EMT. Primary endocardial endothelial cells isolated from embryonic stage 15-18 chick heart explants expressed annexin II protein by Western blot and indirect immunofluorescence. Embryonic quail cells

prepared in an analogous fashion were identified as endocardial by QH1 antibody staining. A spectrofluorometric...

...fold increase in the rate of plasmin generation on chick endocardial cell surfaces versus an acellular control. Plasmin generation was tPA-, plasminogen-, and cell surface-dependent, and was inhibited 90% by...

...heart explants with alpha-2 antiplasmin or anti-annexin II antibody blocked transformation of endocardial cells, as well as their invasion into the collagen matrix, by approx 90% compared to untreated control explants. Western blot analysis of serum-free conditioned medium from heart explant cultures treated with alpha-2 antiplasmin showed a 50% increase in latent TGF-beta3 complex compared with serum-free conditioned medium from untreated explants. This result suggests that inhibition of plasmin activity leads to a...

...of cardiac valves and septae. Annexin II mediated plasmin activity may promote transformation of endocardial cells by activation of latent TGF-beta3, matrix remodeling, and outward migration of the transformed endocardial cells during normal cardiac morphogenesis.

6/3,K/16 (Item 3 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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0001679127 BIOSIS NO.: 19664700083229

A contribution to the study of antibody synthesis. Immunization and activation of spleen ribosomes [Engl. summ.]

ORIGINAL LANGUAGE TITLE: Contribution a l'etude de la biosynthese des anti-corps. Immunisation et activation des ribosomes de la rate [Engl. summ.]

AUTHOR: PAGOULATOS G N

AUTHOR ADDRESS: Lab Biol. Cellulaire, Fac. Sci., Paris, France

JOURNAL: ANN INST PASTEUR 110 ((4)): p497-519 1966 1966

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: Unspecified

...ABSTRACT: microsomal proteins of immunized rabbit spleen, when these microsomes are incubated in presence of soluble acellular fractions of rabbit spleen, or in presence of the same fractions of rat liver. In...

...intense. Incorporation of labelled amino acids into "heavy ribosomes", of a given weight of spleen tissue, incubated in a mixed system with soluble fractions of rat liver, is more intense when...

...One comes therefore to the conclusion that, in immunized animals, a given weight of spleen tissue contains a greater number of active ribosomes than in non-immunized animals. If immunized rabbits...

...between ribosomes activation and antibody synthesis, as well as their common dependance upon the antigenic stimulus, suggest that a synthesized messenger RNA activates the ribosomes and might be related with the...

6/3,K/17 (Item 4 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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0000599166 BIOSIS NO.: 19492300000980

Two unusual sclero-corneal neoplasms

AUTHOR: LOEWENSTEIN A; FOSTER J

JOURNAL: BRIT JOUR OPHTHALMOL 32 ((1)): p1-12 1948 1948

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: Unspecified

...ABSTRACT: element which could not be sharply defined from the tumor.

There were polymorphs, lymphocytes, plasma cells, mast cells and a few giant cells in the periphery of the tumor and in associated tissues. This distinctive picture suggests a tissue reaction to an agent which produces cellular division, and a virus etiology, on the lines...

...of the right cornea in a female of 65 yrs. The tumor was a relatively acellular fibroma with area of calcareous and myxomatous degeneration. The portion of cornea free from tumor was covered by a vascular degenerative pannus. Numerous mast cells were present in the tumor and in the pannus. This suggests an initial chronic inflammation...

...the growth in the pannus. It is postulated that a virus may have been the stimulus which converted the pannus into a neoplasm. ABSTRACT

AUTHORS: K. C. Wybar

6/3,K/18 (Item 1 from file: 73)

DIALOG(R)File 73:EMBASE

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07428753 EMBASE No: 1998337976

Tissue modifications

Black K.S.; Goldstein S.; Ollerenshaw J.

Dr. K.S. Black, CryoLife, Inc, 1655 Roberts Blvd NW, Kennesaw, GA 30144
United States

Transplantation Proceedings (TRANSPLANT. PROC.) (United States) 1998,
30/6 (2729-2731)

CODEN: TRPPA ISSN: 0041-1345

PUBLISHER ITEM IDENTIFIER: S0041134598007982

DOCUMENT TYPE: Journal; Conference Paper

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 28

...number of wide ranging technologies used in experimental models to address the concept of modifying tissue immunogenicity with the goal of improving graft survival. The relationship of these modifications to eventual...

...despite warm culture and cryopreservation, allografted heart valves still are capable of provoking an anti-donor specific immune response in the recipient. Yet there are also no definitive studies that clearly demonstrate in humans that donor-recipient matching improves the durability of these grafts. It seems that some combination of these methodologies for tissue modification used in conjunction with other technological advances will be necessary to yield a new type of graft construct. Our laboratory has been investigating the possibility of repopulating acellular connective tissue matrices with exogenous cells. In this technique appropriate cells obtained from the intended graft recipient are grown into an acellular donor-derived connective tissue matrix. With appropriate graft conditioning it is hoped that these cells can be induced to perform appropriately and serve to enhance

and extend allograft performance. The combination of an acellular allo- or xenogeneic matrix with self cells should provide an optimal graft with very low immunogenicity.

6/3,K/19 (Item 2 from file: 73)
 DIALOG(R)File 73:EMBASE
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04004031 EMBASE No: 1989173027

Molecular basis of fertilization

Garbers D.L.

Howard Hughes Medical Institute, Department of Pharmacology, Vanderbilt University Medical Center, Nashville, TN 37232-0295 United States

Annual Review of Biochemistry (ANNU. REV. BIOCHEM.) (United States) 1989, 58/- (719-742)

CODEN: ARBOA ISSN: 0066-4154

DOCUMENT TYPE: Journal

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

...Here, specific activation of sperm motility, attraction of spermatozoa to the egg, adhesion of sperm cells to the egg, induction of an acrosome reaction, membrane fusion between the gametes, and subsequent...

...of animals, and examples will be discussed in detail later. Eggs are generally enveloped in acellular matrices and/or adherent cells that the spermatozoon must first encounter. In much of the literature, the sites of specific interaction between the spermatozoon and the egg and/or the cellular and acellular components of the egg are not clear for a given species. Therefore, when we speak of molecules associated with the egg, the possible emanance of such molecules from cellular or acellular structures surrounding the egg or from the egg, itself, as well as the deposition of active molecules to the egg-associated structures by cells of the female reproductive tract are all included. It is important to realize that fertilization...

...of relative and absolute specificity cawn be observed with small peptides found in the egg- conditioned media of various animals that interact with cell surface receptors of spermatozoa. In closely related...

...is visible even at the highest peptide concentrations. The biochemical and biological responses of sperm cells to a specific effector molecule are generally similar or identical. Therefore, it is reasonable to...

...and in a receptor molecule (spermatozoa) would occur such that fertilization between the mutant germ cells and spermatozoa or eggs of the general population would be markedly reduced or zero, but...
 ?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECELLULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2	137086	(PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR CONDITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR CELLS OR DONOR)
S3	76	S1 AND S2
S4	0	S3 AND (VECTOR OR TRANSFECTED)
S5	39	RD S3 (unique items)
S6	19	S5 NOT PY>2003

S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
?

S (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRIX))
Processing
Processing
Processing
Processing
Processing
Processing
Processing

 300551 STIMULUS
 34934407 IN
 1339881 VIVO
 1294235 IN(W) VIVO
 571377 EXTRACELLULAR
 481080 MATRIX
 145462 EXTRACELLULAR(W)MATRIX
S8 120 (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W)
 MATRIX))

?

S S1 AND S4
 3991 S1
 0 S4
S9 0 S1 AND S4

?

Set Items Description
S1 3991 (DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECEL-
 LULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2 137086 (PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR COND-
 ITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR
 CELLS OR DONOR)
S3 76 S1 AND S2
S4 0 S3 AND (VECTOR OR TRANSFECTED)
S5 39 RD S3 (unique items)
S6 19 S5 NOT PY>2003
S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S8 120 (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI-
 X))
S9 0 S1 AND S4
?

S S1 AND S8
 3991 S1
 120 S8
S10 0 S1 AND S8

?

Set Items Description
S1 3991 (DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECEL-
 LULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2 137086 (PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR COND-
 ITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR
 CELLS OR DONOR)
S3 76 S1 AND S2
S4 0 S3 AND (VECTOR OR TRANSFECTED)

S5 39 RD S3 (unique items)
 S6 19 S5 NOT PY>2003
 S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 S8 120 (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI-
 X))
 S9 0 S1 AND S4
 S10 0 S1 AND S8
 ?

S (BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
 1380858 BONE
 507422 MARROW
 571377 EXTRACELLULAR
 481080 MATRIX
 S11 65 (BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
 ?

S S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 3991 S1
 3264403 TISSUE
 359561 REPAIR
 325834 ENGINEERING
 37197 TISSUE(W) (REPAIR OR ENGINEERING)
 S12 840 S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 ?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECEL- LULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2	137086	(PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR COND- ITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR CELLS OR DONOR)
S3	76	S1 AND S2
S4	0	S3 AND (VECTOR OR TRANSFECTED)
S5	39	RD S3 (unique items)
S6	19	S5 NOT PY>2003
S7	1	S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S8	120	(STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI- X))
S9	0	S1 AND S4
S10	0	S1 AND S8
S11	65	(BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
S12	840	S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
?		

S S11 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 65 S11
 3264403 TISSUE
 359561 REPAIR
 325834 ENGINEERING
 37197 TISSUE(W) (REPAIR OR ENGINEERING)
 S13 0 S11 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 ?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECEL- LULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2	137086	(PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR COND-


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            ITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR
            CELLS OR DONOR)
S3          76      S1 AND S2
S4          0       S3 AND (VECTOR OR TRANSFECTED)
S5          39      RD S3  (unique items)
S6          19      S5 NOT PY>2003
S7          1       S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S8          120     (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI-
            X))
S9          0       S1 AND S4
S10         0       S1 AND S8
S11         65      (BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
S12         840     S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S13         0       S11 AND (TISSUE (W) (REPAIR OR ENGINEERING))
?

RD S11
    S14        36   RD S11  (unique items)
?

S S14 NOT PY>2003
    36   S14
    6743292 PY>2003
    S15        30   S14 NOT PY>2003
?

S S15 AND (CONTROL AND EXPERIMENTAL)
    30   S15
    5114473 CONTROL
    4731494 EXPERIMENTAL
    S16        1   S15 AND (CONTROL AND EXPERIMENTAL)
?

T S16/3,K/ALL

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16/3,K/1 (Item 1 from file: 155)
 DIALOG(R)File 155:MEDLINE(R)
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12809659 PMID: 10920430

Alterations in proteins of bone marrow extracellular matrix in
 undernourished mice.

Vituri C L; Alvarez-Silva M; Trentin A G; Borelli P
 Departamento de Analises Clinicas, Centro de Ciencias da Saude,
 Universidade Federal de Santa Catarina, Florianopolis, SC, Brasil.

Brazilian journal of medical and biological research = Revista brasileira
 de pesquisas medicas e biologicas / Sociedade Brasileira de Biofisica ...
 et al. (BRAZIL) Aug 2000, 33 (8) p889-95, ISSN 0100-879X--Print

Journal Code: 8112917

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Alterations in proteins of bone marrow extracellular matrix in
 undernourished mice.

... present study was to determine the effect of protein malnutrition on
 the glycoprotein content of bone marrow extracellular matrix (ECM).
 Two-month-old male Swiss mice were submitted to protein malnutrition with a

low-protein diet containing 4% casein as compared to 20% casein in the control diet. When the experimental group had attained a 20% loss of their original body weight, we extracted the ECM...

... samples by SDS-PAGE (7.5%) and ECL Western blotting. Quantitative differences were observed between control and experimental groups. Bone marrow ECM from undernourished mice had greater amounts of extractable fibronectin (1.6-fold increase) and laminin (4.8-fold increase) when compared to the control group. These results suggest an association between fluctuations in the composition of the hematopoietic microenvironment...

; Animals; Blotting, Western; Case- Control Studies; Electrophoresis, Polyacrylamide Gel; Fibronectins--analysis--AN; Hematopoiesis, Extramedullary; Laminin--analysis--AN; Mice
?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECELLULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2	137086	(PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR CONDITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR CELLS OR DONOR)
S3	76	S1 AND S2
S4	0	S3 AND (VECTOR OR TRANSFECTED)
S5	39	RD S3 (unique items)
S6	19	S5 NOT PY>2003
S7	1	S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S8	120	(STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI-X))
S9	0	S1 AND S4
S10	0	S1 AND S8
S11	65	(BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
S12	840	S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S13	0	S11 AND (TISSUE (W) (REPAIR OR ENGINEERING))
S14	36	RD S11 (unique items)
S15	30	S14 NOT PY>2003
S16	1	S15 AND (CONTROL AND EXPERIMENTAL)

?

T S15/3,K/ALL

15/3,K/1 (Item 1 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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14267864 PMID: 12694247

Hyaluronan, a major non-protein glycosaminoglycan component of the extracellular matrix in human bone marrow, mediates dexamethasone resistance in multiple myeloma.

Vincent Thierry; Molina Laurence; Espert Lucile; Mechti Nadir
INSERM Unite U475 and UMR-CNRS5094, Montpellier, and Laboratoire d'Hematologie, Hopital St-Eloi, Montpellier, France.

British journal of haematology (England) Apr 2003, 121 (2) p259-69,
ISSN 0007-1048--Print Journal Code: 0372544

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... activity of different cytokines or growth factors. As HA is a major component of the bone marrow extracellular matrix, these findings support the idea that HA could play a major role in the survival...

15/3,K/2 (Item 2 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

13788515 PMID: 12056556

Structural similarity between the bone marrow extracellular matrix protein and neurokinin 1 could be the limiting factor in the hematopoietic effects of substance P.

Rameshwar Pranela; Gascon Pedro; Bandari Persis S; Joshi Deval D; Fernandes Annemarie; Dang Anju

Department of Medicine, UMDNJ-New Jersey Medical School, Newark 07103, USA. rameshwa@umdnj.edu

Canadian journal of physiology and pharmacology (Canada) May 2002, 80 (5) p475-81, ISSN 0008-4212--Print Journal Code: 0372712

Contract/Grant No.: CA89868; CA; NCI; HL-54973; HL; NHLBI; HL-57675; HL; NHLBI

Publishing Model Print

Document type: Journal Article; Research Support, U.S. Gov't, P.H.S.

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Structural similarity between the bone marrow extracellular matrix protein and neurokinin 1 could be the limiting factor in the hematopoietic effects of substance...

15/3,K/3 (Item 3 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
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12809659 PMID: 10920430

Alterations in proteins of bone marrow extracellular matrix in undernourished mice.

Vituri C L; Alvarez-Silva M; Trentin A G; Borelli P

Departamento de Analises Clinicas, Centro de Ciencias da Saude, Universidade Federal de Santa Catarina, Florianopolis, SC, Brasil.

Brazilian journal of medical and biological research = Revista brasileira de pesquisas medicas e biologicas / Sociedade Brasileira de Biofisica ... et al. (BRAZIL) Aug 2000, 33 (8) p889-95, ISSN 0100-879X--Print

Journal Code: 8112917

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Alterations in proteins of bone marrow extracellular matrix in undernourished mice.

... present study was to determine the effect of protein malnutrition on the glycoprotein content of bone marrow extracellular matrix (ECM). Two-month-old male Swiss mice were submitted to protein malnutrition with a low...

15/3,K/4 (Item 4 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

11265222 PMID: 9058710
CD44-mediated adhesiveness of human hematopoietic progenitors to hyaluronan is modulated by cytokines.
Legras S; Levesque J P; Charrad R; Morimoto K; Le Bousse C; Clay D; Jasmin C; Smadja-Joffe F
Institut National de la Sante et de la Recherche Medicale U268, Hopital Paul Brousse, Villejuif, France.
Blood (UNITED STATES) Mar 15 1997, 89 (6) p1905-14, ISSN 0006-4971
--Print Journal Code: 7603509
Publishing Model Print
Document type: Journal Article; Research Support, Non-U.S. Gov't
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

...mediate adhesion of human CD34+ HPC to immobilized hyaluronan (HA), an abundant glycosaminoglycan of the bone marrow extracellular matrix. Our data show that, although CD34+ cells strongly express CD44, only 13.3% +/- 1.1...

15/3,K/5 (Item 5 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

10613728 PMID: 7556532
Marrow-derived heparan sulfate proteoglycan mediates the adhesion of hematopoietic progenitor cells to cytokines.
Bruno E; Luikart S D; Long M W; Hoffman R
SyStemix, Palo Alto, CA, USA.
Experimental hematology (UNITED STATES) Oct 1995, 23 (11) p1212-7, ISSN 0301-472X--Print Journal Code: 0402313
Contract/Grant No.: CA45279-04; CA; NCI; CA49419; CA; NCI; HL42674-04; HL; NHLBI
Publishing Model Print
Document type: Journal Article; Research Support, U.S. Gov't, P.H.S.
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Heparan sulfate proteoglycan (HS-PG), an important component of the human bone marrow extracellular matrix (ECM), is believed to influence hematopoietic progenitor cell development by binding and localizing growth factors...

15/3,K/6 (Item 6 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

10442947 PMID: 7537448
Ovine bone marrow extracellular matrix and soluble protein extraction: fetuin amino terminus microheterogeneity.
Peters C; Budde C L; Breon T A; Kuper A; Kim J
Department of Pediatrics, University of Iowa College of Medicine, Iowa

City 52242-1083, USA.

American journal of the medical sciences (UNITED STATES) May 1995, 309
(5) p285-94, ISSN 0002-9629--Print Journal Code: 0370506
Contract/Grant No.: P30 HD-27748; HD; NICHD
Publishing Model Print
Document type: Comparative Study; Journal Article; Research Support,
Non-U.S. Gov't; Research Support, U.S. Gov't, P.H.S.
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Ovine bone marrow extracellular matrix and soluble protein
extraction: fetuin amino terminus microheterogeneity.

15/3,K/7 (Item 7 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

10214348 PMID: 7957660
Effects of glycosaminoglycans on U-937 leukemia cell proliferation and
differentiation: structure-function relationship.
Volpi N; Petrini M; Conte A; Valentini P; Venturelli T; Bolognani L;
Ronca G
Department of Biologia Animale, University of Modena, Italy.
Experimental cell research (UNITED STATES) Nov 1994, 215 (1) p119-30
, ISSN 0014-4827--Print Journal Code: 0373226
Publishing Model Print
Document type: Comparative Study; Journal Article
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

... in order to clarify the control of development and differentiation of
hematopoietic progenitor cells by bone marrow extracellular matrix .
Heparin from beef intestinal mucosa, heparan sulfate from beef spleen,
dermatan sulfate from beef intestinal...

15/3,K/8 (Item 8 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

10086584 PMID: 8014614
Effects of extracellular matrix proteins on macrophage differentiation,
growth, and function: comparison of liquid and agar culture systems.
Armstrong J W; Chapes S K Spooner B S KS St U, Manhattan
Division of Biology, NASA Specialized Center of Research and Training,
Kansas State University, Manhattan 66506.
Journal of experimental zoology (UNITED STATES) Jul 1 1994, 269 (3)
p178-87, ISSN 0022-104X--Print Journal Code: 0375365
Publishing Model Print
Document type: Journal Article; Research Support, U.S. Gov't, Non-P.H.S.
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

... The mechanism behind this reduction in haematopoiesis has yet to be
elucidated. However, changes in bone marrow extracellular matrix
(ECM) may be involved. To further understand the role of ECM products in

macrophage differentiation...

15/3,K/9 (Item 9 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

10027340 PMID: 8163649

Bone marrow extracellular matrix molecules improve gene transfer into human hematopoietic cells via retroviral vectors.

Moritz T; Patel V P; Williams D A

Herman B Wells Center for Pediatric Research, James Whitcomb Riley Hospital for Children, Indianapolis, Indiana 46202-5225.

Journal of clinical investigation (UNITED STATES) Apr 1994, 93 (4)
p1451-7, ISSN 0021-9738--Print Journal Code: 7802877

Contract/Grant No.: P01 HL-45168; HL; NHLBI; R01 HL-46528; HL; NHLBI

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't; Research Support, U.S. Gov't, P.H.S.

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Bone marrow extracellular matrix molecules improve gene transfer into human hematopoietic cells via retroviral vectors.

... target cells compared with infection with viral supernatant. We have investigated the role of defined bone marrow extracellular matrix molecules (ECM) in this phenomenon. Here we report that infection of cells adhering to the...

15/3,K/10 (Item 10 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

09648846 PMID: 7682944

Haemonectin, a granulocytic-cell-binding protein, is related to the plasma glycoprotein fetuin.

White H; Totty N; Panayotou G

Department of Oncology, University College and Middlesex School of Medicine, London, England.

European journal of biochemistry / FEBS (GERMANY) Apr 1 1993, 213 (1)
p523-8, ISSN 0014-2956--Print Journal Code: 0107600

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Haemonectin, a protein present in rabbit bone marrow extracellular matrix extracts, has been reported to bind granulocytes in a developmentally regulated manner. We have purified...

15/3,K/11 (Item 11 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

09354120 PMID: 1504379

Bone marrow matrix promotes differentiation and prolongs the cell cycle

of U-937 cells.

Hamdan H F; Luikart S D
Department of Medicine, University of Minnesota Medical School,
Minneapolis.

Oncology research (UNITED STATES) 1992, 4 (4-5) p201-7, ISSN
0965-0407--Print Journal Code: 9208097

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... growth and differentiation of a variety of cell types. In this study,
the effects of bone marrow extracellular matrix on U-937 cells, a
human histiocytic lymphoma cell line, were assessed. Sixty percent of...

... matrix (2980 cpm/10(6) cells vs 230 cpm/10(6) cells on plastic).
Furthermore, bone marrow extracellular matrix inhibited
proliferation of U-937 cells. After four days in culture, there was a 65...
... 5 hr prolongation in cycle length in cells grown on extracellular
matrix. We conclude that bone marrow extracellular matrix induced
macrophage-like differentiation and inhibited proliferation of U-937 cells
with a prolongation of...

15/3,K/12 (Item 12 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2007 Dialog. All rts. reserv.

09184948 PMID: 1845236

Immunohistochemical and ultrastructural study of human bone marrow
extracellular matrix.

Lucena S B; Cotta-Pereira G

UERJ/UFRJ, Departamento de Histologia, Brasil.

Memorias do Instituto Oswaldo Cruz (BRAZIL) 1991, 86 Suppl 3 p115,
ISSN 0074-0276--Print Journal Code: 7502619

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Immunohistochemical and ultrastructural study of human bone marrow
extracellular matrix .

15/3,K/13 (Item 13 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2007 Dialog. All rts. reserv.

09140358 PMID: 1809376

Bone marrow extracellular matrix induces HL-60 cells to produce an
autonomous differentiation factor.

Mane S; Winkelmann J C; Luikart S D

Department of Medicine, University of Minnesota Medical School,
Minneapolis 55455.

Cell growth & differentiation - the molecular biology journal of the
American Association for Cancer Research (UNITED STATES) Dec 1991, 2
(12) p637-43, ISSN 1044-9523--Print Journal Code: 9100024

Publishing Model Print

Document type: Journal Article; Research Support, Non-U.S. Gov't
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Bone marrow extracellular matrix induces HL-60 cells to produce
an autonomous differentiation factor.

15/3,K/14 (Item 14 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

08385770 PMID: 2403825

Fetal expression of hemonectin: an extracellular matrix hematopoietic
cytoadhesion molecule.

Peters C; O'Shea K S; Campbell A D; Wicha M S; Long M W
Department of Pediatrics, University of Michigan Medical School, Ann
Arbor.

Blood (UNITED STATES) Jan 15 1990, 75 (2) p357-64, ISSN 0006-4971--
Print Journal Code: 7603509

Contract/Grant No.: HL 07622; HL; NHLBI; HL 35255; HL; NHLBI
Publishing Model Print

Document type: Journal Article; Research Support, U.S. Gov't, P.H.S.
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Hemonectin, a component of bone marrow extracellular matrix, is a
lineage- and organ-specific attachment molecule for cells of the
granulocytic lineage. We...

15/3,K/15 (Item 15 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2007 Dialog. All rts. reserv.

08020774 PMID: 2783573

Reproducible establishment of hemopoietic supportive stromal cell lines
from murine bone marrow.

Itoh K; Tezuka H; Sakoda H; Konno M; Nagata K; Uchiyama T; Uchino H; Mori
K J

Department of Biology, Faculty of Science, Niigata University, Japan.
Experimental hematology (UNITED STATES) Feb 1989, 17 (2) p145-53,
ISSN 0301-472X--Print Journal Code: 0402313

Publishing Model Print
Document type: Journal Article
Languages: ENGLISH
Main Citation Owner: NLM
Record type: MEDLINE; Completed

Descriptors: *Bone Marrow ; * Extracellular Matrix --physiology--PH;
*Hematopoiesis; *Hematopoietic Stem Cells--physiology--PH

15/3,K/16 (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

17781780 BIOSIS NO.: 200400148441

Hyaluronan, an essential component of the hematopoietic niche, facilitates recovery of ablated hematopoiesis by regulation of cytokine production.
AUTHOR: Khaldoyanidi Sophia K (Reprint); Matrosova Vera Y; Orlovskaya Irina A; Serobyann Naira (Reprint); McClelland Michael
AUTHOR ADDRESS: Department of Vascular Biology, La Jolla Institute for Molecular Medicine, San Diego, CA, USA**USA
JOURNAL: Blood 102 (11): p839a November 16, 2003 2003
MEDIUM: print
CONFERENCE/MEETING: 45th Annual Meeting of the American Society of Hematology San Diego, CA, USA December 06-09, 2003; 20031206
SPONSOR: American Society of Hematology
ISSN: 0006-4971
DOCUMENT TYPE: Meeting; Meeting Poster; Meeting Abstract
RECORD TYPE: Abstract
LANGUAGE: English

...ABSTRACT: HA), a member of glycosaminoglycan family is present in bone marrow, where it participates in bone marrow extracellular matrix (ECM) assembly. We have recently demonstrated that HA is not a passive structural element of...

15/3,K/17 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

17781242 BIOSIS NO.: 200400147903
Impact of VLA-5 expression on the reconstitution properties of hematopoietic stem cells mobilized by cyclophosphamide/G-CSF.
AUTHOR: Wierenga Pieter K (Reprint); de Haan Gerald (Reprint); Weersing Ellen (Reprint); van Os Ronald (Reprint)
AUTHOR ADDRESS: Department of Stem Cell Biology, University of Groningen, Groningen, Netherlands**Netherlands
JOURNAL: Blood 102 (11): p697a November 16, 2003 2003
MEDIUM: print
CONFERENCE/MEETING: 45th Annual Meeting of the American Society of Hematology San Diego, CA, USA December 06-09, 2003; 20031206
SPONSOR: American Society of Hematology
ISSN: 0006-4971
DOCUMENT TYPE: Meeting; Meeting Poster; Meeting Abstract
RECORD TYPE: Abstract
LANGUAGE: English

...ABSTRACT: Since VLA-5 has been implicated in the adhesive interactions of stem cells with the bone marrow extracellular matrix and stromal cells, we unexpectedly found an inverse relationship between hematopoietic reconstitution and the percentage...

15/3,K/18 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

17745833 BIOSIS NO.: 200400116590
Transplantation of Liv-8 negative fraction of bone marrow cells reverses CCl4-induced liver fibrosis.
AUTHOR: Sakaida Isao (Reprint); Aoyama Koji (Reprint); Yamamoto Naoki (Reprint); Ishikawa Tsuyoshi (Reprint); Omori Kaoru (Reprint); Terai Shuji (Reprint); Nishina Hiroshi; Okita Kiwamu (Reprint)
AUTHOR ADDRESS: Yamaguchi University, Ube, Yamaguchi, Japan**Japan

JOURNAL: Hepatology 38 (4 Suppl. 1): p223A October 2003 2003
MEDIUM: print
CONFERENCE/MEETING: 54th Annual Meeting of the American Association for the Study of Liver Diseases Boston, MA, USA October 24-28, 2003; 20031024
SPONSOR: American Association for the Study of Liver Diseases
ISSN: 0270-9139 (ISSN print)
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Abstract
LANGUAGE: English

DESCRIPTORS:

...ORGANISMS: PARTS ETC: bone marrow extracellular matrix --

15/3,K/19 (Item 4 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

17208125 BIOSIS NO.: 200300166844
Adhesion to bone marrow extracellular matrix proteins modulates osteogenic differentiation of human mesenchymal stem cells.
AUTHOR: Salaszyk R M (Reprint); Batorsky A (Reprint); Plopper G E (Reprint)
AUTHOR ADDRESS: Department of Biology, Rensselaer Polytechnic Institute, Troy, NY, USA**USA
JOURNAL: Molecular Biology of the Cell 13 (Supplement): p346a Nov. 2002 2002
MEDIUM: print
CONFERENCE/MEETING: 42nd Annual Meeting of the American Society for Cell Biology San Francisco, CA, USA December 14-18, 2002; 20021214
SPONSOR: American Society for Cell Biology
ISSN: 1059-1524
DOCUMENT TYPE: Meeting; Meeting Abstract
RECORD TYPE: Citation
LANGUAGE: English

Adhesion to bone marrow extracellular matrix proteins modulates osteogenic differentiation of human mesenchymal stem cells.
DESCRIPTORS:
CHEMICALS & BIOCHEMICALS: bone marrow extracellular matrix proteins...

15/3,K/20 (Item 5 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rts. reserv.

15168189 BIOSIS NO.: 199900427849
Comparison of bone marrow extracellular matrices
AUTHOR: Lee Myeongwoo (Reprint); Christopherson Indu P; Lehman Jeffrey M; Bennett Cory J; Cheung H Tak
AUTHOR ADDRESS: Department of Molecular of Molecular Biology, Princeton University, Princeton, NJ, 08544, USA**USA
JOURNAL: Biochimica et Biophysica Acta 1428 (2-3): p300-304 Aug. 5, 1999 1999
MEDIUM: print
ISSN: 0006-3002
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

DESCRIPTORS:

ORGANISMS: PARTS ETC: adult bovine bone marrow extracellular matrix --...

...fetal bovine bone marrow extracellular matrix --

15/3,K/21 (Item 6 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

(c) 2007 The Thomson Corporation. All rts. reserv.

14673034 BIOSIS NO.: 199800467281

Murine bone marrow extracellular matrix study by confocal laser scanning microscopy

AUTHOR: Pelajo-Machado M; Mota E M; Vale L S; Lenzi J A; Lenzi H L

AUTHOR ADDRESS: Dep. Pathol., Inst. Oswaldo Cruz, Fundacao Oswaldo Cruz, Rio de Janeiro, RJ, Brazil**Brazil

JOURNAL: Experimental Hematology (Charlottesville) 26 (8): p704 Aug., 1998 1998

MEDIUM: print

CONFERENCE/MEETING: 27th Annual Meeting of the International Society for Experimental Hematology Vancouver, British Columbia, Canada August 1-5, 1998; 19980801

SPONSOR: International Society for Experimental Hematology

ISSN: 0301-472X

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Citation

LANGUAGE: English

Murine bone marrow extracellular matrix study by confocal laser scanning microscopy

DESCRIPTORS:

ORGANISMS: PARTS ETC: bone marrow extracellular matrix

15/3,K/22 (Item 7 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

(c) 2007 The Thomson Corporation. All rts. reserv.

14099694 BIOSIS NO.: 199799733754

Plastic embedded undecalcified bone biopsies: An immunohistochemical method for routine study of bone marrow extracellular matrix

AUTHOR: Lucena S B (Reprint); Duarte M E L; Fonseca E C

AUTHOR ADDRESS: Hosp. Universitario Antonio Pedro, Dep. Patologia, Rua Marques de Parana 303, Niteroi, RJ 24030, Brazil**Brazil

JOURNAL: Journal of Histotechnology 20 (3): p253-257 1997 1997

ISSN: 0147-8885

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

Plastic embedded undecalcified bone biopsies: An immunohistochemical method for routine study of bone marrow extracellular matrix

DESCRIPTORS:

MISCELLANEOUS TERMS: ... BONE MARROW EXTRACELLULAR MATRIX ;

15/3,K/23 (Item 8 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

(c) 2007 The Thomson Corporation. All rts. reserv.

10430899 BIOSIS NO.: 199140073790

ADHESIVE PROPERTIES OF HUMAN ERYTHROBLASTIC PRECURSOR CELLS

AUTHOR: COULOMBEL L (Reprint); VUILLET-GAUGLER M H; LEROY C; ROSEMBLATT M;
BRETON-GORIUS J

AUTHOR ADDRESS: INSTITUTE DE PATHOLOGIE CELLULAIRE, HOPITAL DE BICETRE,
94270 KREMLIN-BICETRE, FRANCE**FRANCE

JOURNAL: Blood Cells (New York) 17 (1): p65-82 1991

CONFERENCE/MEETING: SYMPOSIUM ON ERYTHROBLASTIC ISLANDS, PARIS, FRANCE,
NOVEMBER 1989. BLOOD CELLS (N Y).

ISSN: 0340-4684

DOCUMENT TYPE: Meeting

RECORD TYPE: Citation

LANGUAGE: ENGLISH

DESCRIPTORS: HUMAN HEMATOPOIETIC DIFFERENTIATION BONE MARROW
EXTRACELLULAR MATRIX FIBRONECTIN

15/3,K/24 (Item 9 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

(c) 2007 The Thomson Corporation. All rts. reserv.

09334175 BIOSIS NO.: 198936043066

**A HEPARIN-HEPARAN SULFATE PROTEOGLYCAN-ASSOCIATED FRACTION OF BONE MARROW
EXTRACELLULAR MATRIX CAN INDUCE MATURATION OF HL-60 MYELOID LEUKEMIA
CELLS**

AUTHOR: LUIKART S D (Reprint); FURCHT L T; MCCARTHY J B; OEGEMA T R JR

AUTHOR ADDRESS: DEP MED, UNIV MINNESOTA, MINNEAPOLIS, MINN, USA**USA

JOURNAL: Clinical Research 36 (6): p853A 1988

CONFERENCE/MEETING: ABSTRACTS SUBMITTED TO THE ANNUAL MEETING OF THE
MIDWEST SECTION, AMERICAN FEDERATION FOR CLINICAL RESEARCH, CHICAGO,
ILLINOIS, USA, NOVEMBER 9-11, 1988. CLIN RES.

ISSN: 0009-9279

DOCUMENT TYPE: Meeting

RECORD TYPE: Citation

LANGUAGE: ENGLISH

**A HEPARIN-HEPARAN SULFATE PROTEOGLYCAN-ASSOCIATED FRACTION OF BONE
MARROW EXTRACELLULAR MATRIX CAN INDUCE MATURATION OF HL-60 MYELOID
LEUKEMIA CELLS**

15/3,K/25 (Item 10 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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09242312 BIOSIS NO.: 198886082233

**BCNU-INDUCED INCREASE IN SULFATED GLYCOSAMINOGLYCAN PRODUCTION BY HUMAN
BONE MARROW STROMAL CELLS**

AUTHOR: OGLE K M (Reprint); LUIKART S D

AUTHOR ADDRESS: BOX 325, UNIV MINN HOSP, MINNEAPOLIS, MINN 55455, USA**USA

JOURNAL: Experimental Hematology (Charlottesville) 16 (7): p636-640 1988

ISSN: 0301-472X

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: ENGLISH

DESCRIPTORS: 1 3 BIS-2-CHLOROETHYL-1-NITROSOUREA HEMATOPOIESIS BONE

MARROW EXTRACELLULAR MATRIX MYELOTOXICITY LEUKEMIA

15/3,K/26 (Item 1 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 2007 Elsevier B.V. All rts. reserv.

11266126 EMBASE No: 2001262749
Runt-related gene 2 in endothelial cells: Inducible expression and
specific regulation of cell migration and invasion
Sun L.; Vitolo M.; Passaniti A.
A. Passaniti, University of Maryland, Greenebaum Cancer Center, BRB Room
7-021, 655 West Baltimore Street, Baltimore, MD 21201 United States
AUTHOR EMAIL: apass001@umaryland.edu
Cancer Research (CANCER RES.) (United States) 01 JUL 2001, 61/13
(4994-5001)
CODEN: CNREA ISSN: 0008-5472
DOCUMENT TYPE: Journal ; Article
LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH
NUMBER OF REFERENCES: 55

MEDICAL DESCRIPTORS:

gene expression; angiogenesis; tumor growth; metastasis; cell migration;
cell invasion; osteoblast; lymphoma; bone marrow ; extracellular
matrix ; protein expression; human; human cell; article; nucleotide
sequence; priority journal

15/3,K/27 (Item 2 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 2007 Elsevier B.V. All rts. reserv.

04858608 EMBASE No: 1991353344
Immunochemical localization of extracellular materials in bone marrow of
rats
Hamilton R.; Campbell F.R.
Dept Anatomical Sci/Neurobiol, Health Sciences Center, University of
Louisville, Louisville, KY 40292 United States
Anatomical Record (ANAT. REC.) (United States) 1991, 231/2 (218-224)
CODEN: ANREA ISSN: 0003-276X
DOCUMENT TYPE: Journal; Article
LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

MEDICAL DESCRIPTORS:

* bone marrow ; * extracellular matrix

15/3,K/28 (Item 3 from file: 73)
DIALOG(R)File 73:EMBASE
(c) 2007 Elsevier B.V. All rts. reserv.

04043111 EMBASE No: 1989212153
Bone marrow microenvironment
Minguell J.J.; Fernandez M.; Tetas M.; Martinez J.; Bruzzone M.;
Rodriguez J.P.
Unidad de Biologia Celular, INTA y Departamento de Ciencias Basicas,
Facultad de Medicine, Sede Oriente, Universidad de Chile, Santiago 11
Chile
Archivos de Biologia y Medicina Experimentales (ARCH. BIOL. MED. EXP.)
(Chile) 1988, 21/1 (177-182)

CODEN: ABMXA ISSN: 0004-0533
 DOCUMENT TYPE: Journal
 LANGUAGE: SPANISH SUMMARY LANGUAGE: ENGLISH

MEDICAL DESCRIPTORS:

*acute lymphocytic leukemia; * bone marrow ; * extracellular matrix ; * fibroblast

15/3,K/29 (Item 4 from file: 73)
 DIALOG(R)File 73:EMBASE
 (c) 2007 Elsevier B.V. All rts. reserv.

03770076 EMBASE No: 1988219512

Extracellular matrix of the marrow microenvironment

Gordon M.Y.

Leukaemia Research Fund Centre, Institute of Cancer Research, London, SW3
 United Kingdom

British Journal of Haematology (BR. J. HAEMATOL.) (United Kingdom)
 1988, 70/1 (1-4)

CODEN: BJHEA ISSN: 0007-1048

DOCUMENT TYPE: Journal

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

MEDICAL DESCRIPTORS:

* bone marrow ; * extracellular matrix ; *hematopoiesis; *leukemia
 --etiology--et

15/3,K/30 (Item 5 from file: 73)
 DIALOG(R)File 73:EMBASE
 (c) 2007 Elsevier B.V. All rts. reserv.

03466556 EMBASE No: 1987219137

Haemonectin, a bone marrow adhesion protein specific for cells of granulocyte lineage

Campbell A.D.; Long M.W.; Wicha M.S.

Division of Hematology and Oncology, Department of Internal Medicine,
 University of Michigan, Simpson Research Institute, Ann Arbor, MI 48109

United States

Nature (NATURE) (United Kingdom) 1987, 329/6141 (744-746)

CODEN: NATUA ISSN: 0028-0836

DOCUMENT TYPE: Journal

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

MEDICAL DESCRIPTORS:

* bone marrow ; * extracellular matrix ; *granulocyte; *hematopoietic cell
 ?

Set	Items	Description
S1	3991	(DECELLULARIZED OR ACELLULAR OR DECELLULARISATION OR DECELLULARIZATION) (S) (TISSUE OR MATRIX OR MATRICES)
S2	137086	(PRECONDITIONING OR CONDITIONING OR PRECONDITIONED OR CONDITIONED OR STIMULUS) (S) ((IN (W) VIVO) OR TISSUE OR ORGAN OR CELLS OR DONOR)
S3	76	S1 AND S2
S4	0	S3 AND (VECTOR OR TRANSFECTED)
S5	39	RD S3 (unique items)

S6 19 S5 NOT PY>2003
 S7 1 S6 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 S8 120 (STIMULUS) (S) ((IN (W) VIVO) AND (EXTRACELLULAR (W) MATRI-
 X))
 S9 0 S1 AND S4
 S10 0 S1 AND S8
 S11 65 (BONE (W) MARROW (W) EXTRACELLULAR (W) MATRIX)
 S12 840 S1 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 S13 0 S11 AND (TISSUE (W) (REPAIR OR ENGINEERING))
 S14 36 RD S11 (unique items)
 S15 30 S14 NOT PY>2003
 S16 1 S15 AND (CONTROL AND EXPERIMENTAL)
 ?

COST

02nov07 15:17:49 User259876 Session D1048.2
 \$25.12 7.388 DialUnits File155
 \$6.60 30 Type(s) in Format 3
 \$6.60 30 Types
 \$31.72 Estimated cost File155
 \$49.13 8.188 DialUnits File5
 \$32.20 14 Type(s) in Format 3
 \$32.20 14 Types
 \$81.33 Estimated cost File5
 \$72.89 6.125 DialUnits File73
 \$23.10 7 Type(s) in Format 3
 \$23.10 7 Types
 \$95.99 Estimated cost File73
 OneSearch, 3 files, 21.701 DialUnits FileOS
 \$6.66 INTERNET
 \$215.70 Estimated cost this search
 \$216.73 Estimated total session cost 21.980 DialUnits

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